BORON BY CURCUMIN METHOD SM 4500-B B-2000 (21 st Edition) ADDITIONAL QC REQUIREMENTS FOR THIS METHOD: Certified or Accredited laboratories using this method are assessed to applicable requirements of SM 1020 and SM 4020.						
Facility Name:	VELAP ID					
Assessor Name:Analyst Name:	Analyst Name:		Inspection Date			
Relevant Aspect of Standards	Method Reference	Y	N	N/A	Comments	
Records Examined: SOP Number/ Revision/ Date		Analyst:				
Sample ID: Date of Sample Prepare	Date of Sample Preparation:		Date of Analysis:			
Were samples collected in Polyethylene, Fluoropolymer, or Quartz containers?	40 CFR 136 Table 1I					
Were Non-Potable Water samples acidified to a pH < 2 with HNO3?	40 CFR 136 Table 1I					
Were Non-Potable Water samples held for no longer than 6 months?	40 CFR 136 Table 1I					
Was the filter photometer or spectrophotometer used at 540 nm, and did it have a 1 cm light path?	SM 4500-B B-2000 2.a					
Were all reagents stored in polyethylene or boron-free containers?	SM 4500-B B-2000 3					
Was the Anhydrous Boric Acid kept tightly stoppered to prevent entrance of atmospheric moisture?	SM 4500-B B-2000 3.a					
Was the stock Boron Solution composed of 571.6 mg Anhydrous Boric Acid in 1000 mL distilled water?	SM 4500-B B-2000 3.a					
Was the Curcumin Reagent composed of 40 mg finely ground Curcumin+ 5.0 g oxalic acid+ 4.2 mL concentrated HCl in 100 mL 95% Ethyl Alcohol (OR 100 mL 95% Isopropanol)?	SM 4500-B B-2000 3.c					
Was the Curcumin Reagent stored in the refrigerator for not longer than several days?	SM 4500-B B-2000 3.c					
Were the working calibration standards prepared by adding 4.0 mL of Curcumin Reagent to 1.0 mL of standards followed by allowing the standards to float in a water bath at 55 ± 2°C for 80 minutes?	SM 4500-B B-2000 4.b					
Notes/Comments:						

SM 4500-B B-2000 (21st Edition) **BORON BY CURCUMIN METHOD** ADDITIONAL QC REQUIREMENTS FOR THIS METHOD: Certified or Accredited laboratories using this method are assessed to applicable requirements of SM 1020 and SM 4020. Υ Ν N/A **Comments Relevant Aspect of Standards** Method Reference After the water bath, were standards allowed to cool to room temperature followed by the addition of 10 mL 95% SM 4500-B Ethyl Alcohol (or Isopropyl Alcohol) while stirring until the B-2000 4.b red color completely dissoluted? Were samples treated in the same way as calibration SM 4500-B standards in 4.b? B-2000 4.c Were photometric readings taken within 1 hour of drying SM 4500-B the samples? B-2000 4.b Were samples containing greater than 1.00 mg B/L SM 4500-B diluted with distilled water prior to drying? B-2000 4.c If final sample solutions were turbid, were the sample SM 4500-B solutions filtered through filter paper (Whatman no. 30 or B-2000 4.c equivalent)? If low-level samples (50µg to 200 µg) were analyzed using a visual estimation by visual comparison against 2003 NELAC calibration standards, were the results reported as 5.5.10.3.2.f estimated? (SM 4500-B B-2000 4.d) If high hardness or cation interference was an issue, were samples pipetted into a column containing a SM 4500-B strongly-acidic cation exchange resin at a rate of about 2 B-2000 4.e drops/second prior to treatment? Notes/Comments: